

GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: May 23, 2005, 23:05:16 ; Search time 1868 Seconds
(without alignments)
427.917 Million cell updates/sec

Title: US-10-646-436-10

Perfect score: 21

Sequence: 1 gcagcagagucuccauctt 21

Scoring table: IDENTITY NUC

Gapop 10.0 , Gapext 1.0

Searched: 34239544 seqs, 19032134700 residues

Total number of hits satisfying chosen parameters: 68479088

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : EST:*

- 1: gb_est1:*
- 2: gb_est2:*
- 3: gb_hic:*
- 4: gb_est3:*
- 5: gb_est4:*
- 6: gb_est5:*
- 7: gb_est6:*
- 8: gb_gss1:*
- 9: gb_gss2:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
C 1	19.4	92.4	532	9 CE099065	tigr-gss-
C 2	19.4	92.4	573	2 BF921577	MR1-NT017
C 3	19.0	90.5	85	2 AW276802	XP66a01.x
C 4	19.0	90.5	100	2 BF920141	MR1-NT017
C 5	19.0	90.5	102	4 BF958934	PM1-NN120
C 6	19.0	90.5	119	2 AW901233	CM4-NN101
C 7	19.0	90.5	124	4 BM821731	K-EST0090
C 8	19.0	90.5	136	6 CD612966	56043322J
C 9	19.0	90.5	137	6 CD612965	56043322H
C 10	19.0	90.5	138	5 BQ339862	PM1-NN120
C 11	19.0	90.5	142	7 D45267	HUMHG1194.H
C 12	19.0	90.5	147	5 BQ339466	PM1-NN120
C 13	19.0	90.5	148	2 BF946357	PM1-EN006
C 14	19.0	90.5	149	7 R47195	CBS-389.Sub
C 15	19.0	90.5	169	2 BE766895	RC2-NT011
C 16	19.0	90.5	177	1 AL048592	DKF2P469B
C 17	19.0	90.5	183	4 B1032792	MR4-NN018
C 18	19.0	90.5	184	4 BF958930	PM1-NN120
C 19	19.0	90.5	185	4 B1036860	MR4-NT014
C 20	19.0	90.5	196	1 A1745406	wc37d01.x
C 21	19.0	90.5	197	1 B1036862	MR4-NT014
C 22	19.0	90.5	201	4 BF957666	PM1-NN120
C 23	19.0	90.5	203	4 BF957653	PM1-NN120
C 24	19.0	90.5	207	4 BF957858	PM1-NN120

25	19	90.5	211	2 BF928116	MR4-NT014
C 26	19	90.5	224	2 AW161224	au70a10.y
C 27	19	90.5	226	2 BF935119	MR4-NT014
C 28	19	90.5	236	4 BF948789	MR3-NN021
C 29	19	90.5	238	2 BF923639	MR4-NT014
C 30	19	90.5	239	4 BG898974	HOA21-1-C
C 31	19	90.5	241	1 CR767137	DKF2P469B
C 32	19	90.5	243	1 AA336628	EST41242
C 33	19	90.5	244	2 BE766870	RC2-NT011
C 34	19	90.5	248	2 BF923643	MR4-NT014
C 35	19	90.5	249	4 BF945175	PM1-NN120
C 36	19	90.5	250	5 BQ345410	MR4-NT014
C 37	19	90.5	252	2 BF923633	MR4-NT014
C 38	19	90.5	252	4 BF963107	PM1-NN120
C 39	19	90.5	253	2 BF887875	QV2-TN017
C 40	19	90.5	253	4 B1041998	MR4-NT014
C 41	19	90.5	256	4 B1041248	MR4-NT014
C 42	19	90.5	257	6 CD612964	56037472J
C 43	19	90.5	258	4 BF961150	PM1-NN120
C 44	19	90.5	258	7 CR763018	DKF2P469G
C 45	19	90.5	262	4 BF947155	MR3-NN021

ALIGNMENTS

CE099065 532 bp DNA linear GSS 24-SBP-2003
tigr-gss-dog-17000371093708 Dog Library Canis familiaris genomic,
genomic survey sequence.
CE099065
CE099065.1 GI:35165950
GSS.
Canis familiaris (dog)
Canis familiaris
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Carnivora; Fissipedia; Canidae; Canis.
REFERENCE 1 (bases 1 to 532)
AUTHORS Kirkness,E.F., Bafna,V., Halpern,A.L., Levy,S., Remington,K.,
Rusch,D.B., Delcher,A.L., Pop,M., Wang,W., Fraser,C.M. and
Venter,J.C.
TITLE The dog genome: survey sequencing and comparative analysis
JOURNAL Science 301 (5641), 1898-1903 (2003)
MEDLINE 22875432
PUBMED 14512627
COMMENT Contact: Kirkness EF
The Institute for Genomic Research
Department of Eukaryotic Genomics, TIGR, 9712 Medical Center Drive,
Rockville, MD 20850, USA
Tel: 301-838-0200
Fax: 301-838-0208
Email: ekirkness@tigr.org
Class: shotgun.
Location/Qualifiers
1. 532
/organism="Canis familiaris"
/mol_type="genomic DNA"
/strain="Standard Poodle"
/db_xref="taxon:9615"
/clone_lib="Dog Library"
/note="Site 1: BstXI; Libraries were prepared from
peripheral blood"

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Best Local Similarity 76.2%; Pred. No. 2.1e+02;
Matches 16; Conservative 4; Mismatches 1; Indels 0; Gaps 0;
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Db 82 GCAGCAGAGTCTACATCATTT 62

/clone_lib="NN1011"
 /note="Organ: nervous normal; Vector: puc18; Site_1: SmaI;
 Site_2: SmaI; A mini-library was made by cloning products
 derived from ORESTES PCR (U.S. Letters Patent application
 No. 196,716 - Ludwig Institute for Cancer Research)
 profiles into the pUC 18 vector. Reverse transcription of
 tissue mRNA and cDNA amplification were performed under
 low stringency conditions."

ORIGIN

Query Match 90.5%; Score 19; DB 2; Length 119;
 Best Local Similarity 73.7%; Pred. No. 2.5e+02;
 Matches 14; Conservative 5; Mismatches 0; Indels 0; Gaps 0;

Qy 1 GCAGCAGAGUCUUCACAU 19
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 Db 101 GCAGCAGAGTCTTCATCAT 83

RESULT 7

BM821731/c
 LOCUS BM821731 124 bp mRNA linear EST 06-MAR-2002
 DEFINITION K-EST0090952 S20T665307 Homo sapiens cDNA clone S20T665307-15-H01
 5' mRNA sequence.

ACCESSION BM821731
 VERSION BM821731.1 GI:19178144
 KEYWORDS EST.
 SOURCE Homo sapiens (human)

ORGANISM

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE

AUTHORS Kim,N.S., Hahn,Y., Oh,J.H., Lee,J.Y., Ahn,H.Y., Chu,M.Y., Kim,M.R.,
 Oh,K.J., Cheong,J.E., Sohn,H.Y., Kim,J.M., Park,H.S., Kim,S. and
 Kim,Y.S.

TITLE 21C Frontier Korean EST Project 2001

JOURNAL Unpublished (2002)

COMMENT Contact: Kim YS

Genome Research Center
 Korea Research Institute of Bioscience & Biotechnology
 52 Eoeun-dong Yuseong-gu, Daejeon 305-333, South Korea

Tel: +82-42-860-4470

Fax: +82-42-860-4409

Email: yongsung@mail.kribb.re.kr

Plate: 15 row: H column: 01

High quality sequence stop: 124.

Location/Qualifiers

FEATURES

source

1..124

/organism="Homo sapiens"

/mol_type="mRNA"

/db_xref="taxon:9606"

/clone="S20T665307-15-H01"

/sex="M"

/lab_host="Top10F"

/clone_lib="S20T665307"

/note="Organ: Stomach; Vector: pCNS; Site_1: EcoRI;
 Site_2: NotI; The poly (A)+ RNA was dephosphorylated with
 bacterial alkaline phosphatase (BAP) and then decapped
 with tobacco acid pyrophosphatase (TAP). The decapped
 intact mRNA was ligated with DNA-RNA linker including EcoR
 I site by treatment of T4 RNA ligase and the first strand
 cDNA was synthesized from oligo dt-selected mRNA by
 priming with dt-tailed vector. The dt-tailed vector was
 adjusted to have about 60nt. The cDNA vector was
 circularized with E. coli DNA ligase after digestion of
 EcoRI which site is also included in vector. An RNA strand
 converted to a DNA strand by Okayama-Berg method. The
 obtained cDNA vectors were used for transformation of
 competent cells E. coli Top10F by electroporation method.
 The cDNA libraries constructed by this method are
 full-length enriched cDNA library."

ORIGIN

Query Match 90.5%; Score 19; DB 4; Length 124;
 Best Local Similarity 73.7%; Pred. No. 2.5e+02;
 Matches 14; Conservative 5; Mismatches 0; Indels 0; Gaps 0;

Qy 1 GCAGCAGAGUCUUCACAU 19
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 Db 100 GCAGCAGAGTCTTCATCAT 82

RESULT 8

CD612966
 LOCUS CD612966 136 bp mRNA linear EST 12-JAN-2004
 DEFINITION S6043322J1 FLP Homo sapiens cDNA, mRNA sequence.

ACCESSION CD612966

VERSION CD612966.1 GI:40261230

KEYWORDS EST.

SOURCE Homo sapiens (human)

ORGANISM

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE

1 (bases 1 to 136)

AUTHORS Fu,G.K., Wang,J.T., Yang,J., Au-Young,J. and Stuve,L.L.

TITLE Circular rapid amplification of cDNA ends for high-throughput

JOURNAL extension cloning of partial genes

COMMENT Genomics 84 (1), 205-210 (2004)

Contact: Fu GK

Incyte Genomics, Inc.

3160 Porter Dr., Palo Alto, CA 94304, USA

Tel: 6508454102

Email: gfu@incyte.com.

FEATURES

source

1..136

/organism="Homo sapiens"

/mol_type="mRNA"

/db_xref="taxon:9606"

/clone_lib="FLP"

/note="Vector: pDrive Cloning Vector"

ORIGIN

Query Match 90.5%; Score 19; DB 6; Length 136;
 Best Local Similarity 73.7%; Pred. No. 2.6e+02;
 Matches 14; Conservative 5; Mismatches 0; Indels 0; Gaps 0;

Qy 1 GCAGCAGAGUCUUCACAU 19
 |||||:|:|:|:|:
 Db 71 GCAGCAGAGTCTTCATCAT 89

RESULT 9

CD612965/c
 LOCUS CD612965 137 bp mRNA linear EST 12-JAN-2004
 DEFINITION S6043322H1 FLP Homo sapiens cDNA, mRNA sequence.

ACCESSION CD612965

VERSION CD612965.1 GI:40261229

KEYWORDS EST.

SOURCE Homo sapiens (human)

ORGANISM

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE

1 (bases 1 to 137)

AUTHORS Fu,G.K., Wang,J.T., Yang,J., Au-Young,J. and Stuve,L.L.

TITLE Circular rapid amplification of cDNA ends for high-throughput

JOURNAL extension cloning of partial genes

COMMENT Genomics 84 (1), 205-210 (2004)

Contact: Fu GK

Incyte Genomics, Inc.

3160 Porter Dr., Palo Alto, CA 94304, USA

Tel: 6508454102

Email: gfu@incyte.com.

FEATURES

source

1..137

/organism="Homo sapiens"

/mol_type="mRNA"

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/db_xref="taxon:9606"
/clonelib="FLP"
/note="Vector: pDrive Cloning Vector"

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Best Local Similarity 73.7%; Pred. No. 2.6e+02;
Matches 14; Conservative 5; Mismatches 0; Indels 0; Gaps 0;

QY      1 GCAGCAGAGUCUUAUCAU 19
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Db       66 GCAGCAGAGTCTTCATCAT 48

RESULT 10
BQ339862/c
LOCUS   BQ339862              138 bp mRNA linear EST 20-MAY-2002
DEFINITION PM1-NN1200-011200-010-f07 NN1200 Homo sapiens cDNA, mRNA sequence.
ACCESSION BQ339862
VERSION    BQ339862.1 GI:20999978
KEYWORDS   EST.
SOURCE     Homo sapiens (human)
ORGANISM   Homo sapiens
REFERENCE  1 (bases 1 to 138)
AUTHORS    Dias Neto,E., Garcia Correa,R., Verjovski-Almeida,S., Briones,M.R., Nagai,M.A., da Silva,W. Jr., Zago,M.A., Bordin,S., Costa,F.F., Goldman,G.H., Carvalho,A.P., Matsukuma,A., Baia,G.S., Simpson,D.H., Brunstein,A., deOliveira,P.S., Bucher,P., Jongeneel,C.V., O'Hare,M.J., Soares,F., Brentani,R.R., Reis,L.F., de Souza,S.J. and Simpson,A.J.
TITLE      Shotgun sequencing of the human transcriptome with ORF expressed sequence tags
JOURNAL    Proc. Natl. Acad. Sci. U.S.A. 97 (7), 3491-3496 (2000)
MEDLINE    20202663
COMMENT    10737800
Contact: Simpson A.J.G.
Laboratory of Cancer Genetics
Ludwig Institute for Cancer Research
Rua Prof. Antonio Prudente 109, 4 andar, 01509-010, Sao Paulo-SP, Brazil
Tel.: +55-11-2704922
Fax: +55-11-2707001
Email: asimpson@ludwig.org.br
This sequence was derived from the FAPESP/LICR Human Cancer Genome Project. This entry can be seen in the following URL (http://www.ludwig.org.br/scripts/gethtml.pl?tl=PMI&t2=PM1-NN1200-011200-010-f07&t3=2000-12-01&t4=1)
Seq primer: puc 18 forward
High quality sequence start: 17
High quality sequence stop: 35.
FEATURES             Location/Qualifiers
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                     /clone_lib="NN1200"
                     /note="Organ: nervous normal; Vector: puc18; Site_1: Smal; Site_2: SmaI; A mini-library was made by cloning products derived from ORSTES PCR (U.S. Letters Patent application No. 196,716 - Ludwig Institute for Cancer Research) profiles into the pUC 18 vector. Reverse transcription of tissue mRNA and cDNA amplification were performed under low stringency conditions."

ORIGIN
Query Match          90.5%; Score 19; DB 5; Length 138;
Best Local Similarity 73.7%; Pred. No. 2.6e+02;
Matches 14; Conservative 5; Mismatches 0; Indels 0; Gaps 0;

QY      1 GCAGCAGAGUCUUAUCAU 19
        |||||:::||::||:
Db       66 GCAGCAGAGTCTTCATCAT 48

RESULT 11
D45267/c
LOCUS   D45267              142 bp mRNA linear EST 30-DEC-1995
DEFINITION HUMHG1194 Human cerebral cortex Homo sapiens CDNA, mRNA sequence.
ACCESSION D45267
VERSION    D45267.1 GI:1136645
KEYWORDS   EST.
SOURCE     Homo sapiens (human)
ORGANISM   Homo sapiens
REFERENCE  1 (bases 1 to 142)
AUTHORS    Takahashi,N., Hashida,H., Zhao,N., Misumi,Y. and Sakaki,Y.
TITLE      High-density cDNA filter analysis of the expression profiles of the genes preferentially expressed in human brain
JOURNAL    Gene 164, 219-227 (1995)
MEDLINE    96069586
COMMENT    7590334
Contact: Nobuaki Takahashi
Institute of Medical Science
University of Tokyo
Shirokanedai 4-6-1, Minato-ku, Tokyo, Japan 108
Tel: 03-5449-5625
Fax: 03-5449-5445.
FEATURES             Location/Qualifiers
     source           1..142
                     /organism="Homo sapiens"
                     /mol_type="mRNA"
                     /db_xref="taxon:9606"
                     /clone_lib="Human cerebral cortex"
                     /note="Adult male cerebral cortex tissue."

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Best Local Similarity 73.7%; Pred. No. 2.6e+02;
Matches 14; Conservative 5; Mismatches 0; Indels 0; Gaps 0;

QY      1 GCAGCAGAGUCUUAUCAU 19
        |||||:::||::||:
Db       77 GCAGCAGAGTCTTCATCAT 59

RESULT 12
BQ339466/c
LOCUS   BQ339466              147 bp mRNA linear EST 20-MAY-2002
DEFINITION PM1-NN1200-051100-004-h12 NN1200 Homo sapiens cDNA, mRNA sequence.
ACCESSION BQ339466
VERSION    BQ339466.1 GI:20999152
KEYWORDS   EST.
SOURCE     Homo sapiens (human)
ORGANISM   Homo sapiens
REFERENCE  1 (bases 1 to 147)
AUTHORS    Dias Neto,E., Garcia Correa,R., Verjovski-Almeida,S., Briones,M.R., Nagai,M.A., da Silva,W. Jr., Zago,M.A., Bordin,S., Costa,F.F., Goldman,G.H., Carvalho,A.F., Matsukuma,A., Baia,G.S., Simpson,D.H., Brunstein,A., deOliveira,P.S., Bucher,P., Jongeneel,C.V., O'Hare,M.J., Soares,F., Brentani,R.R., Reis,L.F., de Souza,S.J. and Simpson,A.J.
TITLE      Shotgun sequencing of the human transcriptome with ORF expressed sequence tags
JOURNAL    Proc. Natl. Acad. Sci. U.S.A. 97 (7), 3491-3496 (2000)
MEDLINE    20202663
COMMENT    10737800
Contact: Simpson A.J.G.
Laboratory of Cancer Genetics
Ludwig Institute for Cancer Research
Rua Prof. Antonio Prudente 109, 4 andar, 01509-010, Sao Paulo-SP, Rua Prof. Antonio Prudente 109, 4 andar, 01509-010, Sao Paulo-SP,

```

[illegible]

Brazil
Tel: +55-11-2704922
Fax: +55-11-2707001
Email: asimpson@ludwig.org.br
This sequence was derived from the FAPESP/LICR Human Cancer Genome Project. This entry can be seen in the following URL
(http://www.ludwig.org.br/scripts/gethtml2.pl?l=PM1&t2=PM1-NN1200-051100-004-h12&t3=2000-11-05&t4=1)
Seq primer: puc 18 forward
High quality sequence start: 14
High quality sequence stop: 34.

FEATURES

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/note="Organ: nervous normal; Vector: puc18; Site_1: SmaI; Site_2: SmaI; A mini-library was made by cloning products derived from ORESTES PCR (U.S. Letters Patent application No. 196,716 - Ludwig Institute for Cancer Research) profiles into the pUC 18 vector. Reverse transcription of tissue mRNA and cDNA amplification were performed under low stringency conditions."

ORIGIN

Query Match 90.5%; Score 19; DB 5; Length 147;
Best Local Similarity 73.7%; Pred. No. 2.6e+02;
Matches 14; Conservative 5; Mismatches 0; Indels 0; Gaps 0;

Qy 1 GCAGCAGAGUCUUCAUCAU 19
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Db 124 GCAGCAGAGTCTTCATCAT 142

RESULT 13
BF846357
LOCUS BF846357 148 bp mRNA linear EST 16-JAN-2001
DEFINITION PM1-EN0065-231000-002-b01 EN0065 Homo sapiens cDNA, mRNA sequence.
ACCESSION BF846357
VERSION BF846357.1 GI:12233611
KEYWORDS EST.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 148)
Dias Neto,E., Garcia Correa,R., Verjovski-Almeida,S., Briones,M.R., Nagai,W.A., da Silva,W. Jr., Zago,M.A., Bordin,S., Costa,F.F., Goldman,G.H., Carvalho,A.F., Matsukuma,A., Baia,G.S., Simpson,D.H., Brunstein,A., deoliveira,P.S., Bucher,P., Jongeneel,C.V., O'Hare,M.J., Soares,F., Brentani,R.R., Reis,L.F., de Souza,S.J. and Simpson,A.J.
Shotgun sequencing of the human transcriptome with ORF expressed

sequence tags
Proc. Natl. Acad. Sci. U.S.A. 97 (7), 3491-3496 (2000)
20202663
10737800

FEATURES

source
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/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="CBS-389"
/note="Vector: pBluescript II SK; Site 1: EcoRI; Site 2: XhoI; A subtractive cDNA library was developed by hybridizing antisense, single-stranded phagemid DNA (ssDNA) (as pBluescript SK-) from the ocular ciliary body cDNA library (target) of a 34-year-old female donor in lambda-Uni-ZAP XR with biotinylated sense RNA of an ocular cell line cDNA library (driver) in the same vector."

Contact: Simpson A.J.G.
Laboratory of Cancer Genetics
Ludwig Institute for Cancer Research
Rua Prof. Antonio Prudente 109, 4 andar, 01509-010, Sao Paulo-SP, Brazil
Tel: +55-11-2704922
Fax: +55-11-2707001
Email: asimpson@ludwig.org.br
This sequence was derived from the FAPESP/LICR Human Cancer Genome Project. This entry can be seen in the following URL
(http://www.ludwig.org.br/scripts/gethtml2.pl?l=PM1&t2=PM1-EN0065-231000-002-b01&t3=2000-10-23&t4=1)
Seq primer: puc 18 forward
High quality sequence stop: 110.

FEATURES

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/db_xref="taxon:9606"
/dev_stage="Adult"
/clone_lib="EN0065"
/note="Organ: lung_normal; Vector: puc18; Site_1: SmaI; Site_2: SmaI; A mini-library was made by cloning products derived from ORESTES PCR (U.S. Letters Patent application No. 196,716 - Ludwig Institute for Cancer Research) profiles into the pUC 18 vector. Reverse transcription of tissue mRNA and cDNA amplification were performed under low stringency conditions."

ORIGIN

Query Match 90.5%; Score 19; DB 2; Length 148;
Best Local Similarity 73.7%; Pred. No. 2.6e+02;
Matches 14; Conservative 5; Mismatches 0; Indels 0; Gaps 0;

Qy 1 GCAGCAGUCUUCAUCAU 19
|||||||:|:|:|:
Db 36 GCAGCAGAGTCTTCATCAT 54

RESULT 14

R47195/c
LOCUS R47195 149 bp mRNA linear EST 12-DEC-1995
DEFINITION CBS-389 Subtractive cDNA library ocular ciliary body Homo sapiens cDNA clone CBS-389 5' end similar to TRPM-2 (clusterin) (accession number M64722), mRNA sequence.

ACCESSION R47195
VERSION R47195.1 GI:807537
KEYWORDS EST.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE 1 (bases 1 to 149)
Escribano,J., Ortego,J. and Coca-Prados,M.
Isolation and characterization of cell-specific cDNA clones from a subtractive library of the ocular ciliary body of a single normal human donor: Transcription and synthesis of plasma proteins
J. Biochem. 118 (5), 921-931 (1995)
96318503
MEDLINE 8749308
PUBMED
COMMENT Contact: Coca-Prados, M.
Department of Ophthalmology and Visual Science
Yale University Medical School
330 Cedar Street, New Haven, CT 06520-8061
Tel: 2037852742
Fax: 2037856123
Email: miguel.coca-prados@quickmail.yale.edu
Seq primer: T3.

FEATURES

source
1..149
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="CBS-389"
/note="Vector: pBluescript II SK; Site 1: EcoRI; Site 2: XhoI; A subtractive cDNA library was developed by hybridizing antisense, single-stranded phagemid DNA (ssDNA) (as pBluescript SK-) from the ocular ciliary body cDNA library (target) of a 34-year-old female donor in lambda-Uni-ZAP XR with biotinylated sense RNA of an ocular cell line cDNA library (driver) in the same vector."

ORIGIN

Query Match 90.5%; Score 19; DB 7; Length 149;
Best Local Similarity 73.7%; Pred. No. 2.6e+02;
Matches 14; Conservative 5; Mismatches 0; Indels 0; Gaps 0;

QY 1 GCAGCAGAGUCUUCAU 19
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 Db 69 GCAGCAGAGTCTTCATCAT 51

RESULT 15
 BE766895/c
 LOCUS BE766895 169 bp mRNA linear EST 19-SEP-2000
 DEFINITION RC2-NT0110-050600-013-f03 NT0110 Homo sapiens cDNA, mRNA sequence.
 ACCESSION BE766895
 VERSION BE766895.1 GI:10196819
 KEYWORDS EST.
 SOURCE Homo sapiens (human)

ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 1 (bases 1 to 169)
 Dias Neto, E., Garcia Correa, R., Verjovski-Almeida, S., Briones, M.R.,
 Nagai, M.A., da Silva, W. Jr., Zago, M.A., Bordin, S., Costa, F.F.,
 Goldman, G.H., Carvalho, A.F., Matsukuma, A., Baia, G.S., Simpson, D.H.,
 Brunstein, A., deOliveira, P.S., Bucher, P., Jongeneel, C.V.,
 O'Hare, M.J., Soares, F., Brentani, R.R., Reis, L.F., de Souza, S.J. and
 Simpson, A.J.

TITLE Shotgun sequencing of the human transcriptome with ORF expressed
 sequence tags
 JOURNAL Proc. Natl. Acad. Sci. U.S.A. 97 (7), 3491-3496 (2000)
 MEDLINE 20202663
 PUBMED 10737800
 COMMENT Contact: Simpson A.J.G.
 Laboratory of Cancer Genetics
 Ludwig Institute for Cancer Research
 Rua Prof. Antonio Prudente 109, 4 andar, 01509-010, Sao Paulo-SP,
 Brazil
 Tel: +55-11-2704922
 Fax: +55-11-2707001
 Email: asimpson@ludwig.org.br

FEATURES
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 /mol_type="mRNA"
 /db_xref="taxon:9606"
 /dev_stage="Adult"
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 Site 2: SmaI; A mini-library was made by cloning products
 derived from ORESTES PCR (U.S. Letters Patent application
 No. 196,716 - Ludwig Institute for Cancer Research)
 profiles into the pUC 18 vector. Reverse transcription of
 tissue mRNA and cDNA amplification were performed under
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 High quality sequence start: 18
 High quality sequence stop: 169.

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 Matches 14; Conservative 5; Mismatches 0; Indels 0; Gaps 0;

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Search completed: May 24, 2005, 00:21:54
 Job time : 1874 secs

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C 4	19	90.5	195	2	US-08-485-657A-14	Sequence 14, Appl
C 5	19	90.5	195	4	US-09-366-380-14	Sequence 14, Appl
C 6	19	90.5	195	5	PCT-US95-02303-14	Sequence 14, Appl
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C 8	19	90.5	396	4	US-09-640-173-15	Sequence 15, Appl
C 9	19	90.5	396	4	US-09-713-550-15	Sequence 15, Appl
C 10	19	90.5	396	4	US-09-825-294-15	Sequence 15, Appl
C 11	19	90.5	396	4	US-09-970-966-15	Sequence 15, Appl
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C 13	19	90.5	491	4	US-09-513-998C-3749	Sequence 3749, Ap
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C 15	19	90.5	1648	3	US-09-659-791A-3	Sequence 3, Appl
C 16	19	90.5	1651	3	US-09-659-791A-3	Sequence 13, Appl
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C 23	16.8	80.0	601	4	US-09-949-016-139518	Sequence 139518, A
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C 25	16.8	80.0	117001	4	US-09-949-016-15684	Sequence 15684, A
C 26	16.4	78.1	183112	4	US-09-949-016-14184	Sequence 14184, A
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Sequence 19, Application US/09366380		Sequence 14, Application US/08485657A	
Patent No. 6541603		Patent No. 5942389	
GENERAL INFORMATION:		GENERAL INFORMATION:	
APPLICANT: Kirschling, Deborah J		APPLICANT: Kirschling, Deborah J	
APPLICANT: Gudkov, Andrei		APPLICANT: Gudkov, Andrei	
APPLICANT: Roninson, Igor B		APPLICANT: Roninson, Igor B	
TITLE OF INVENTION: Genes and Genetic Elements Associated		TITLE OF INVENTION: Genes and Genetic Elements Associated	
TITLE OF INVENTION: with Sensitivity to Cisplatin		TITLE OF INVENTION: with Sensitivity to Cisplatin	
NUMBER OF SEQUENCES: 23		NUMBER OF SEQUENCES: 23	
CORRESPONDENCE ADDRESS:		CORRESPONDENCE ADDRESS:	
ADDRESSEE: McDonnell Boehnen Hulbert & Berghoff		ADDRESSEE: McDonnell Boehnen Hulbert & Berghoff	
STREET: 300 South Wacker Drive, 32nd Floor		STREET: 300 South Wacker Drive, 32nd Floor	
CITY: Chicago		CITY: Chicago	
STATE: Illinois		STATE: Illinois	
COUNTRY: USA		COUNTRY: USA	
ZIP: 60606		ZIP: 60606	
COMPUTER READABLE FORM:		COMPUTER READABLE FORM:	
MEDIUM TYPE: Floppy disk		MEDIUM TYPE: Floppy disk	
COMPUTER: IBM PC compatible		COMPUTER: IBM PC compatible	
OPERATING SYSTEM: PC-DOS/MS-DOS		OPERATING SYSTEM: PC-DOS/MS-DOS	
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FILING DATE:		FILING DATE:	
ATTORNEY/AGENT INFORMATION:		ATTORNEY/AGENT INFORMATION:	
NAME: No. 6541603nan, Kevin E		NAME: No. 5942389nan, Kevin E	
REGISTRATION NUMBER: 35,303		REGISTRATION NUMBER: 35,303	
REFERENCE/DOCKET NUMBER: 93,354-N		REFERENCE/DOCKET NUMBER: 93,354-N	
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GENERAL INFORMATION:		GENERAL INFORMATION:	
APPLICANT:		APPLICANT: Kirschling, Deborah J	
TITLE OF INVENTION: Genes and Genetic Elements Associated		TITLE OF INVENTION: Genes and Genetic Elements Associated	
TITLE OF INVENTION: with Sensitivity to Cisplatin		TITLE OF INVENTION: with Sensitivity to Cisplatin	
NUMBER OF SEQUENCES: 25		NUMBER OF SEQUENCES: 23	
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COMPUTER: IBM PC compatible		COMPUTER: IBM PC compatible	
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FILING DATE:		FILING DATE: 07-JUN-1995	
CLASSIFICATION:		CLASSIFICATION: 800	
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APPLICATION NUMBER: 08/199,900		APPLICATION NUMBER: 08/199,900	
FILING DATE:		FILING DATE:	
ATTORNEY/AGENT INFORMATION:		ATTORNEY/AGENT INFORMATION:	
NAME: No. 6541603nan, Kevin E		NAME: No. 5942389nan, Kevin E	
REGISTRATION NUMBER: 35,303		REGISTRATION NUMBER: 35,303	
REFERENCE/DOCKET NUMBER: 93,354-N		REFERENCE/DOCKET NUMBER: 93,354-N	
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; GENERAL INFORMATION:
; APPLICANT: Kirschling, Deborah J
; APPLICANT: Gudkov, Andrei
; APPLICANT: Roninson, Igor B
; TITLE OF INVENTION: Genes and Genetic Elements Associated
; TITLE OF INVENTION: with Sensitivity to Cisplatin
; NUMBER OF SEQUENCES: 23
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: McDonnell Boehnen Hulbert & Berghoff
; STREET: 300 South Wacker Drive, 32nd Floor
; CITY: Chicago
; STATE: Illinois
; COUNTRY: USA
; ZIP: 60606
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
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; OPERATING SYSTEM: PC-DOS/MS-DOS
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; FILING DATE:
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; APPLICATION NUMBER: 08/199,900
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; ATTORNEY/AGENT INFORMATION:
; NAME: No. 6541603nan, Kevin E
; REGISTRATION NUMBER: 35,303
; REFERENCE/DOCKET NUMBER: 93,354-N
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; TELEPHONE: 312-913-0001
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; TITLE OF INVENTION: Genes and Genetic Elements Associated
; TITLE OF INVENTION: with Sensitivity to Cisplatin
; NUMBER OF SEQUENCES: 25
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; Patent No. 6783961
; GENERAL INFORMATION:
; APPLICANT: Dumas Milne Edwards, J.B.
; APPLICANT: Duclert, A. Y.
; APPLICANT: Giordano, J.Y.
; TITLE OF INVENTION: Expressed Sequence Tags and Encoded Human Proteins.
; Patent No. 6783961
; FILE REFERENCE: 59.US2.REG
; CURRENT APPLICATION NUMBER: US/09/513,999C
; CURRENT FILING DATE: 2000-02-24
; PRIOR APPLICATION NUMBER: US 60/122,487
; PRIOR FILING DATE: 1999-02-26
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; Patent No. 6613515
; GENERAL INFORMATION:
; APPLICANT: Xu, Jiangchun
; APPLICANT: Stolk, John A.
; TITLE OF INVENTION: OVARIAN TUMOR SEQUENCES AND
; TITLE OF INVENTION: METHODS OF USE THEREFOR
; FILE REFERENCE: 210121.484C2
; CURRENT APPLICATION NUMBER: US/09/640,173
; CURRENT FILING DATE: 2000-08-15
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; SEQ ID NO 15
; LENGTH: 396
; TYPE: DNA
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RESULT 9

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; Sequence 15, Application US/09713550
; Patent No. 6617109
; GENERAL INFORMATION:
; APPLICANT: Xu, Jiangchun
; APPLICANT: Stolk, John A.
; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR THE
; FILE REFERENCE: 210121.484C4
; CURRENT APPLICATION NUMBER: US/09/713,550
; CURRENT FILING DATE: 2000-11-14
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US-09-713-550-15

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; GENERAL INFORMATION:
; APPLICANT: Xu, Jiangchun
; APPLICANT: Stolk, John A.
; APPLICANT: Algate, Paul A.
; APPLICANT: Fling, Steven P.
; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR THE
; FILE REFERENCE: 210121.484C5
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; CURRENT FILING DATE: 2001-04-03
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; OTHER INFORMATION: n = A,T,C or G
US-09-825-294-15

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Db 77 GCAGCAGAGTCTTCATCAT 59

Matches 14; Conservative 5; Mismatches 0; Indels 0; Gaps 0;

RESULT 11
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; Sequence 15, Application US/09970966
; Patent No. 6720146
; GENERAL INFORMATION:
; APPLICANT: Stolk, John A.
; APPLICANT: Molesh, David Alan
; APPLICANT: Fling, Steven P.
; APPLICANT: Xu, Jiangchun
; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR THE THERAPY
; FILE REFERENCE: 210121.484C6
; CURRENT APPLICATION NUMBER: US/09/970,966
; CURRENT FILING DATE: 2001-10-02
; NUMBER OF SEQ ID NOS: 215
; SOFTWARE: FastSeq for Windows Version 4.0
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US-09-970-966-15

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Db 77 GCAGCAGAGTCTTCATCAT 59

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US-09-621-976-13267/c
; Sequence 13267, Application US/09621976
; Patent No. 6639063
; GENERAL INFORMATION:
; APPLICANT: Dumas Milne Edwards, J.B.
; APPLICANT: Jobert, S.
; APPLICANT: Giordano, J.Y.
; TITLE OF INVENTION: ESTs and Encoded Human Proteins.
; FILE REFERENCE: GENSET.054PR2
; CURRENT APPLICATION NUMBER: US/09/621,976
; CURRENT FILING DATE: 2000-07-21
; NUMBER OF SEQ ID NOS: 19335
; SOFTWARE: Patent.pm
; SEQ ID NO 13267
; LENGTH: 482
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-621-976-13267

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Best Local Similarity 73.7%; Pred. No. 24;
Matches 14; Conservative 5; Mismatches 0; Indels 0; Gaps 0;

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Db 94 GCAGCAGAGTCTTCATCAT 76

RESULT 13

US-09-513-999C-3749/c
; Sequence 3749, Application US/09513999C
; Patent No. 6783961
; GENERAL INFORMATION:
; APPLICANT: Dumas Milne Edwards, J.B.
; APPLICANT: Duclert, A.

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; APPLICANT: Giordano, J.Y.
; TITLE OF INVENTION: Expressed Sequence Tags and Encoded Human Proteins.
; Patent No. 6783961
; FILE REFERENCE: 59.US2.REG
; CURRENT APPLICATION NUMBER: US/09/513,999C
; CURRENT FILING DATE: 2000-02-24
; PRIOR APPLICATION NUMBER: US 60/122,487
; PRIOR FILING DATE: 1999-02-26
; NUMBER OF SEQ ID NOS: 36681
; SOFTWARE: Patent.pm
; SEQ ID NO 3749
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; ORGANISM: Homo sapiens
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; NAME/KEY: CDS
; LOCATION: 108..455
; FEATURE:
; NAME/KEY: sig_peptide
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; OTHER INFORMATION: seq LLFVGLLTWESG/QV
; FEATURE:
; NAME/KEY: misc_feature
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; US-09-513-999C-3749

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Query Match	90.5%	Score 19;	DB 4;	Length 491;
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Db	126	GCACGAGAGTCTTCAT	TCAT	108

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; Sequence 201815, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
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; ORGANISM: Human
US-09-949-016-201815

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RESULT 15
US-09-659-791A-3/C

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; Sequence 3, Application US/09659791A
; Patent No. 6383808
; GENERAL INFORMATION:
; APPLICANT: Brett P. Monia
; APPLICANT: Susan M. Freier
; TITLE OF INVENTION: ANTISENSE MODULATION OF CLUSTERIN EXPRESSION
; FILE REFERENCE: RYS-0156
; CURRENT APPLICATION NUMBER: US/09/659,791A
; CURRENT FILING DATE: 2000-09-11
; NUMBER OF SEQ ID NOS: 90
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GenCore version 5.1.6
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Run on: May 23, 2005, 21:40:56 ; Search time 255 Seconds
(without alignments)
487.508 Million cell updates/sec

Title: US-10-646-436-10

Perfect score: 21

Sequence: 1 gcagcagacucuaucatt 21

Scoring table: IDENTITY_NUC

Gapop 10.0 , Gapext 1.0

Searched: 4390206 seqs, 2959870667 residues

Total number of hits satisfying chosen parameters: 8780412

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

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2: Geneseqn1990s:*
3: Geneseqn2000s:*
4: Geneseqn2001as:*
5: Geneseqn2001bs:*
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12: Geneseqn2004as:*
13: Geneseqn2004bs:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	21	100.0	21	12	ADL70465
2	21	100.0	21	12	ADL70431
3	19	90.5	19	12	ADL70522
4	19	90.5	19	12	ADL70523
5	19	90.5	19	12	ADL70444
6	19	90.5	19	12	ADL70445
7	19	90.5	21	3	AAA94226
8	19	90.5	21	10	ACF36398
9	19	90.5	21	11	ADM83069
10	19	90.5	21	12	ADL70464
11	19	90.5	21	12	ADL70430
12	19	90.5	21	12	ADL70406
13	19	90.5	23	12	ADL70521
14	19	90.5	195	2	AAT00416
15	19	90.5	275	3	AAC06940
16	19	90.5	396	4	AAF94824
17	19	90.5	396	6	ABL48774
18	19	90.5	396	6	ABT03091
19	19	90.5	396	11	ADM10684
20	19	90.5	396	12	ADJ11014

C 21	19	90.5	396	12	ADM43275	Adm43275 Human ova
C 22	19	90.5	461	9	ACH44960	Ach44960 Human foe
C 23	19	90.5	462	9	ACH15312	Ach15312 Human adu
C 24	19	90.5	465	9	ACH44965	Ach44965 Human foe
C 25	19	90.5	490	9	ACH25299	Ach25299 Human adu
C 26	19	90.5	491	3	AAC03751	Aac03751 Human tes
C 27	19	90.5	491	9	ACH30243	Ach30243 Human tes
C 28	19	90.5	492	9	ACH43944	Ach43944 Human foe
C 29	19	90.5	512	2	AAV89150	Aav89150 EST clone
C 30	19	90.5	572	2	AZ42136	Az42136 Human nor
C 31	19	90.5	704	6	ABQ56105	Abq56105 Human ova
C 32	19	90.5	922	3	AAA43857	Aaa43857 Human sec
C 33	19	90.5	1024	10	ABZ83527	Abz83527 Toxicolog
C 34	19	90.5	1067	8	ACC90611	Acc90611 Human CGD
C 35	19	90.5	1117	8	ACC90613	Acc90613 Human CGD
C 36	19	90.5	1315	8	ACC90621	Acc90621 Human CGD
C 37	19	90.5	1369	8	ACC90610	Acc90610 Human CGD
C 38	19	90.5	1373	8	ACC90622	Acc90622 Human CGD
C 39	19	90.5	1451	10	ADI02673	Adi02673 Human cDN
C 40	19	90.5	1568	5	AA344948	Aas44948 cDNA enco
C 41	19	90.5	1589	6	ABS78654	Abs78654 Human cDN
C 42	19	90.5	1610	8	ACC90609	Acc90609 Human CGD
C 43	19	90.5	1648	6	ABN99656	Abn99656 Human clu
C 44	19	90.5	1651	2	AAQ11503	Aaq11503 Cytolysis
C 45	19	90.5	1651	6	ABN99666	Abn99666 Human clu

ALIGNMENTS

RESULT 1

ADL70465

ID ADL70465 standard; RNA; 21 BP.

XX AC ADL70465;

XX DT 20-MAY-2004 (first entry)

XX DE RNAi for human clusterin.

XX KW RNA interference; RNAi; short interfering RNA; siRNA; human; clusterin;

XX KW cytostatic; neuroprotective; nontropic; gene silencing; DNA-RNA hybrid;

XX KW ss.

XX OS Homo sapiens.

XX OS Synthetic.

XX FH Key Location/Qualifiers

XX FT modified_base 20..21

XX FT /*tag= a

XX FT /mod_base= OTHER

XX FT /note= "OTHER= dTdT"

XX PN WO2004018676-A2.

XX PD 04-MAR-2004.

XX PF 21-AUG-2003; 2003WO-CA001277.

XX PR 21-AUG-2002; 2002US-0405193P.

XX PR 03-SEP-2002; 2002US-0408152P.

XX PR 20-MAY-2003; 2003US-0472387P.

XX PA (UYBR-) UNIV BRITISH COLUMBIA.

XX PI Jansen B, Gleave ME, Signaevsky M, Beraldi E, Trougakos IP;

XX PI Gonos ES;

XX PI WPI; 2004-226852/21.

XX DR New RNA molecule less than 49 bases and having a sequence effective to

XX PT mediate degradation or block translation of mRNA that is the

XX PT transcriptional product of a target gene, useful for treating Alzheimer's

XX PT

```

PT disease or cancer.
XX
PS Claim 4; SEQ ID NO 10; 63pp; English.
XX
CC The present sequence is the antisense strand of a short interfering RNA
CC (siRNA) targeted to human clusterin. The sense strand is also provided
CC ADL70464. The siRNA can be used to interfere with the expression of
CC clusterin. Clusterin, also known as testosterone-repressed prostate
CC message-2 (TRPM-2) or sulfated glycoprotein-2 (SGP-2), is expressed in
CC increased amounts by prostate tumour cells following androgen withdrawal,
CC and has also been shown to be critical for neuritic toxicity in mouse
CC models of Alzheimer's disease. siRNAs of the invention can be used alone
CC or in combination with other chemotherapy or apoptosis inducing
CC treatments for the treatment of prostate cancer, sarcomas such as
CC osteosarcoma, renal cell carcinoma, breast cancer, bladder cancer, lung
CC cancer, colon cancer, ovarian cancer, anaplastic large cell lymphoma and
CC melanoma, and also for the treatment of Alzheimer's disease.
XX
SQ Sequence 21 BP; 5 A; 5 C; 4 G; 2 T; 5 U; 0 Other;

Query Match      100.0%; Score 21; DB 12; Length 21;
Best Local Similarity 100.0%; Pred. No. 3.8;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GCAGCAGAGUCUUCUCAUUTT 21
   |||||
DB 1 GCAGCAGAGUCUUCUCAUUTT 21

RESULT 2
ADL70431
ID ADL70431 standard; RNA; 21 BP.
XX
AC ADL70431;
XX
DT 20-MAY-2004 (first entry)
XX
DE RNAi for human clusterin.
XX
KW Human; clusterin; RNAi; melanoma; cytostatic; gene silencing;
KW short interfering RNA; siRNA; DNA-RNA hybrid; ss.
XX
OS Homo sapiens.
OS Synthetic.
XX
FH Key Location/Qualifiers
FT modified_base 20..21
FT /*tag= a
FT /mod_base= OTHER
FT /note= "Tt"
XX
PN WO2004018675-A1.
XX
PD 04-MAR-2004.
XX
PF 21-AUG-2003; 2003WO-CA001276.
XX
PR 21-AUG-2002; 2002US-0405193P.
PR 03-SEP-2002; 2002US-0408152P.
PR 02-DEC-2002; 2002US-0319748P.
PR 20-MAY-2003; 2003US-0472387P.
XX
PA (UYBR-) UNIV BRITISH COLUMBIA.
PA (GLEA/) GLEAVE M E.
XX
PI Jansen B;
XX
DR WPI; 2004-226851/21.
XX
PT Treating melanoma in a mammalian subject comprises administering to the
PT subject a therapeutic agent effective to reduce the effective amount of
PT clusterin in the melanoma cells.
XX

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PS Claim 20; SEQ ID NO 29; 32pp; English.
XX
CC The present sequence is that of a short interfering RNA (siRNA) molecule
CC targeted to human clusterin ADL70403. The invention relates to the
CC treatment of melanoma through reduction in the effective amount of
CC clusterin. The therapeutic agent may be an antisense oligonucleotide
CC ADL70404-ADL70421 or short interfering RNA (siRNA) ADL70422-ADL70445
CC targeted to clusterin. The siRNAs molecules direct cleavage of clusterin
CC mRNA. A method for regulating expression of bcl-xL in a subject or cell
CC line comprises administering an agent effective to modulate the amount of
CC clusterin expression. In clusterin-expressing cells, expression of bcl-xL
CC is down-regulated when the effective amount of clusterin is reduced. Such
CC inhibition is significant because bcl-xL is known to act as an inhibitor
CC of apoptosis.
XX
SQ Sequence 21 BP; 5 A; 5 C; 4 G; 2 T; 5 U; 0 Other;

Query Match      100.0%; Score 21; DB 12; Length 21;
Best Local Similarity 100.0%; Pred. No. 3.8;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GCAGCAGAGUCUUCUCAUUTT 21
   |||||
DB 1 GCAGCAGAGUCUUCUCAUUTT 21

RESULT 3
ADL70522/c
ID ADL70522 standard; RNA; 19 BP.
XX
AC ADL70522;
XX
DT 20-MAY-2004 (first entry)
XX
DE RNAi for human clusterin.
XX
KW RNA interference; RNAi; short interfering RNA; siRNA; human; clusterin;
KW cytostatic; neuroprotective; nontropic; gene silencing; DNA-RNA hybrid;
KW ss.
XX
OS Homo sapiens.
OS Synthetic.
XX
FH Key Location/Qualifiers
FT modified_base 18..19
FT /*tag= a
FT /mod_base= OTHER
FT /note= "OTHER= dtdt"
XX
PN WO2004018676-A2.
XX
PD 04-MAR-2004.
XX
PF 21-AUG-2003; 2003WO-CA001277.
XX
PR 21-AUG-2002; 2002US-0405193P.
PR 03-SEP-2002; 2002US-0408152P.
PR 20-MAY-2003; 2003US-0472387P.
XX
PA (UYBR-) UNIV BRITISH COLUMBIA.
XX
PI Jansen B, Gleave ME, Signaevsky M, Beraldi E, Trougakos IP;
PI Gonos ES;
XX
DR WPI; 2004-226852/21.
XX
PT New RNA molecule less than 49 bases and having a sequence effective to
PT mediate degradation or block translation of mRNA that is the
PT transcriptional product of a target gene, useful for treating Alzheimer's
PT disease or cancer.
XX
PS Claim 4; SEQ ID NO 67; 63pp; English.
XX

```


CC The present sequence is the sense strand of a short interfering RNA
CC (siRNA) targeted to a specific portion ADL70521 of human clusterin cDNA.
CC The antisense strand is also provided ADL70523. The siRNA can be used to
CC interfere with the expression of clusterin. Clusterin, also known as
CC testosterone-repressed prostate message-2 (TRPM-2) or sulfated
CC glycoprotein-2 (SGP-2), is expressed in increased amounts by prostate
CC tumour cells following androgen withdrawal, and has also been shown to be
CC critical for neuritic toxicity in mouse models of Alzheimer's disease.
CC siRNAs of the invention can be used alone or in combination with other
CC chemotherapies or apoptosis inducing treatments for the treatment of
CC prostate cancer, sarcomas such as osteosarcoma, renal cell carcinoma,
CC breast cancer, bladder cancer, lung cancer, colon cancer, ovarian cancer,
CC anaplastic large cell lymphoma and melanoma, and also for the treatment
CC of Alzheimer's disease. In an example from the invention, the present
CC siRNA was used to examine the effects of clusterin gene silencing in PC-3
CC prostate cancer cells. A reduction in clusterin transcript was observed.
XX
SQ Sequence 19 BP; 5 A; 4 C; 5 G; 0 T; 5 U; 0 Other;

Query Match 90.5%; Score 19; DB 12; Length 19;
Best Local Similarity 73.7%; Pred. No. 33;
Matches 14; Conservative 5; Mismatches 0; Indels 0; Gaps 0;
QY 1 GCAGCAGAGUCUUCUACAU 19
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DB 19 GCAGCAGAGTCTTCATCAT 1

RESULT 4
ADL70523
ID ADL70523 standard; RNA; 19 BP.
XX
AC ADL70523;
XX
DT 20-MAY-2004 (first entry)
XX
DE RNAi for human clusterin.
XX
KW RNA interference; RNAi; short interfering RNA; siRNA; human; clusterin;
KW cytostatic; neuroprotective; neutropic; gene silencing; DNA-RNA hybrid;
KW ss.
XX Homo sapiens.
OS Synthetic.
XX
FH Key Location/Qualifiers
FT modified_base 18..19
FT /*tag= a
FT /mod_base= OTHER
FT /note= "OTHER= dtdt"
XX
PN WO2004018676-A2.
XX
XX 04-MAR-2004.
XX
XX 21-AUG-2003; 2003WO-CA001277.
XX
PR 21-AUG-2002; 2002US-0405193P.
PR 03-SEP-2002; 2002US-0408152P.
PR 20-MAY-2003; 2003US-0472387P.
XX
PA (UYBR-) UNIV BRITISH COLUMBIA.
XX
XX Jansen B, Gleave ME, Signaevsky M, Beraldi E, Trogakos IP;
PI Gonos ES;
XX
XX WPI; 2004-226852/21.
XX
XX New RNA molecule less than 49 bases and having a sequence effective to
PT mediate degradation or block translation of mRNA that is the
PT transcriptional product of a target gene, useful for treating Alzheimer's
PT disease or cancer.
XX

PS Claim 4; SEQ ID NO 68; 63pp; English.
XX
CC The present sequence is the antisense strand of a short interfering RNA
CC (siRNA) targeted to a specific portion ADL70521 of human clusterin cDNA.
CC The sense strand is also provided ADL70522. The siRNA can be used to
CC interfere with the expression of clusterin. Clusterin, also known as
CC testosterone-repressed prostate message-2 (TRPM-2) or sulfated
CC glycoprotein-2 (SGP-2), is expressed in increased amounts by prostate
CC tumour cells following androgen withdrawal, and has also been shown to be
CC critical for neuritic toxicity in mouse models of Alzheimer's disease.
CC siRNAs of the invention can be used alone or in combination with other
CC chemotherapies or apoptosis inducing treatments for the treatment of
CC prostate cancer, sarcomas such as osteosarcoma, renal cell carcinoma,
CC breast cancer, bladder cancer, lung cancer, colon cancer, ovarian cancer,
CC anaplastic large cell lymphoma and melanoma, and also for the treatment
CC of Alzheimer's disease. In an example from the invention, the present
CC siRNA was used to examine the effects of clusterin gene silencing in PC-3
CC prostate cancer cells. A reduction in clusterin transcript was observed.
XX
SQ Sequence 19 BP; 5 A; 5 C; 4 G; 0 T; 5 U; 0 Other;
Query Match 90.5%; Score 19; DB 12; Length 19;
Best Local Similarity 100.0%; Pred. No. 33;
Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 GCAGCAGAGUCUUCUACAU 19
|||||||:|:|:|:
DB 1 GCAGCAGAGUCUUCUACAU 19
XX
RESULT 5
ADL70444/C
ID ADL70444 standard; RNA; 19 BP.
XX
AC ADL70444;
XX
DT 20-MAY-2004 (first entry)
XX
DE RNAi for human clusterin.
XX
KW Human; clusterin; RNAi; melanoma; cytostatic; gene silencing;
KW short interfering RNA; siRNA; DNA-RNA hybrid; ss.
XX
XX Homo sapiens.
OS Synthetic.
XX
FH Key Location/Qualifiers
FT modified_base 18..19
FT /*tag= a
FT /mod_base= OTHER
FT /note= "OTHER= TT"
XX
PN WO2004018675-A1.
XX
XX 04-MAR-2004.
XX
XX 21-AUG-2003; 2003WO-CA001276.
XX
PR 21-AUG-2002; 2002US-0405193P.
PR 03-SEP-2002; 2002US-0408152P.
PR 02-DEC-2002; 2002US-0319748P.
PR 20-MAY-2003; 2003US-0472387P.
XX
XX (UYBR-) UNIV BRITISH COLUMBIA.
PA
PA (GLEA/) GLEAVE M E.
XX
XX Jansen B;
XX
XX WPI; 2004-226851/21.
XX
XX Treating melanoma in a mammalian subject comprises administering to the
XX subject a therapeutic agent effective to reduce the effective amount of
XX clusterin in the melanoma cells.
PT

XX PS Claim 20; SEQ ID NO 42; 32pp; English.

XX CC The present sequence is that of a short interfering RNA (siRNA) molecule

XX CC targeted to human clusterin ADL70403. The invention relates to the

XX CC treatment of melanoma through reduction in the effective amount of

XX CC clusterin. The therapeutic agent may be an antisense oligonucleotide

XX CC targeted to clusterin. The siRNAs molecules direct cleavage of clusterin

XX CC ADL70404-ADL70421 or short interfering RNA (siRNA) ADL70422-ADL70445

XX CC mRNA. A method for regulating expression of bcl-xL in a subject or cell

XX CC line comprises administering an agent effective to modulate the amount of

XX CC clusterin expression. In clusterin-expressing cells, expression of bcl-xL

XX CC is down-regulated when the effective amount of clusterin is reduced. Such

XX CC inhibition is significant because bcl-xL is known to act as an inhibitor

XX CC of apoptosis.

XX SQ Sequence 19 BP; 5 A; 4 C; 5 G; 0 T; 5 U; 0 Other;

Query Match 90.5%; Score 19; DB 12; Length 19;

Best Local Similarity 73.7%; Pred. No. 33;

Matches 14; Conservative 5; Mismatches 0; Indels 0; Gaps 0;

Qy 1 GCAGCAGAGUCUUCUCAU 19

Db 19 GCAGCAGAGTCTTCATCAT 1

RESULT 6

ID ADL70445 standard; RNA; 19 BP.

XX AC ADL70445;

XX DT 20-MAY-2004 (first entry)

XX DE RNAi for human clusterin.

XX KW Human; clusterin; RNAi; melanoma; cytostatic; gene silencing;

XX KW short interfering RNA; siRNA; DNA-RNA hybrid; ss.

XX OS Homo sapiens.

XX OS Synthetic.

XX FH Key Location/Qualifiers

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FT /*tag= a

-FT /mod_base= OTHER

FT /*note= "OTHER= TT"

XX WO2004018675-A1.

XX PD 04-MAR-2004.

XX PF 21-AUG-2003; 2003WO-CA001276.

XX PR 21-AUG-2002; 2002US-0405193P.

PR 03-SEP-2002; 2002US-0408152P.

PR 02-DEC-2002; 2002US-0319748P.

PR 20-MAY-2003; 2003US-0472387P.

XX (UYBR-) UNIV BRITISH COLUMBIA.

PA (GLEA/) GLEAVE M E.

XX Jansen B;

XX WPI; 2004-226851/21.

XX Treating melanoma in a mammalian subject comprises administering to the

PT subject a therapeutic agent effective to reduce the effective amount of

PT clusterin in the melanoma cells.

XX Claim 20; SEQ ID NO 43; 32pp; English.

CC The present sequence is that of a short interfering RNA (siRNA) molecule

CC targeted to human clusterin ADL70403. The invention relates to the

CC treatment of melanoma through reduction in the effective amount of

CC clusterin. The therapeutic agent may be an antisense oligonucleotide

CC ADL70404-ADL70421 or short interfering RNA (siRNA) ADL70422-ADL70445

CC targeted to clusterin. The siRNAs molecules direct cleavage of clusterin

CC mRNA. A method for regulating expression of bcl-xL in a subject or cell

CC line comprises administering an agent effective to modulate the amount of

CC clusterin expression. In clusterin-expressing cells, expression of bcl-xL

CC is down-regulated when the effective amount of clusterin is reduced. Such

CC inhibition is significant because bcl-xL is known to act as an inhibitor

CC of apoptosis.

XX SQ Sequence 19 BP; 5 A; 5 C; 4 G; 0 T; 5 U; 0 Other;

Query Match 90.5%; Score 19; DB 12; Length 19;

Best Local Similarity 100.0%; Pred. No. 33;

Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 GCAGCAGAGUCUUCUCAU 19

Db 1 GCAGCAGAGUCUUCUCAU 19

RESULT 7

AAA94226

ID AAA94226 standard; DNA; 21 BP.

XX AC AAA94226;

XX DT 12-JAN-2001 (first entry)

XX DE Human testosterone-repressed prostate message-2 antisense oligo #2.

XX KW Human; testosterone-repressed prostate message-2; TRPM-2; clusterin;

XX KW sulfated glycoprotein-2; SGP-2; cancer; antisense oligonucleotide; ss.

XX OS Homo sapiens.

XX PN WO200049937-A2.

XX PD 31-AUG-2000.

XX PF 25-FEB-2000; 2000WO-US004875.

PR 26-FEB-1999; 99US-0121726P.

XX (UYBR-) UNIV BRITISH COLUMBIA.

XX Gleave M, Rennie PS, Miyake H, Nelson C;

XX WPI; 2000-533132/48.

XX Treating prostatic tumors and renal cancers by antisense inhibition of

PT the testosterone-repressed prostate messenger-2 gene.

XX Claim 3; Page 36; 38pp; English.

XX The present sequence is an antisense oligonucleotide directed at the

CC human testosterone-repressed prostate message-2 (TRPM-2, also known as

CC clusterin, sulfated glycoprotein-2 or SGP-2). The sequence was shown to

CC promote the regression of tumours, and oligonucleotides directed at human

CC TRPM-2 can be used in the treatment of tumour cells expressing the TRPM-2

CC gene. These include prostate cancer, renal cell cancer and some breast

CC cancer cells. In addition to this, they also increase the

CC chemosensitivity of the cells, meaning that conventional chemotherapy is

CC more effective

XX SQ Sequence 21 BP; 6 A; 6 C; 4 G; 5 T; 0 U; 0 Other;

Query Match 90.5%; Score 19; DB 3; Length 21;

Best Local Similarity 73.7%; Pred. No. 34;

Matches 14; Conservative 5; Mismatches 0; Indels 0; Gaps 0;

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 Db 3 GCAGCAGAGTCTTCATCAT 21

RESULT 8

ACF36398
 ID ACF36398 standard; DNA; 21 BP.

AC ACF36398;
 XX

DT 18-DEC-2003 (first entry)

XX TRPM-2 antisense oligonucleotide.
 XX

KW TRPM-2; testosterone-repressed prostate message-2; cytostatic; androgen;
 KW prostate cancer; anti-apoptotic protein; antisense; ss.

XX Synthetic.

OS Homo sapiens.

XX WO2003072591-A1.
 PN

XX 04-SEP-2003.
 PD

XX 20-FEB-2003; 2003WO-US005305.
 PF

XX 22-FEB-2002; 2002US-00080794.
 PR

XX (UYBR-) UNIV BRITISH COLUMBIA.
 PA

XX Gleave M, Rennie PS, Miyake H, Nelson C, Monia BP;
 PI

XX WPI; 2003-689981/65.
 DR

XX New modified antisense oligonucleotide, useful particularly for treating
 PT prostatic cancer, inhibits the testosterone-repressed prostate message-2.
 PT

XX Claim 1; Page 25; 4pp; English.
 PS

XX The invention relates to a compound consisting of an oligonucleotide with
 CC a phosphorothioate backbone throughout, in which: (a) sugars on
 CC nucleotide residues 1-4 and 18-21 are 2'-O-methoxyethyl modified, and the
 CC remaining nucleotides 5-17 are 2'-deoxy; and (b) the cytosines at
 CC positions 1, 4 and 19 are 5-methylated. Oligonucleotide shown in sequence
 CC ACF36398 (I) is used: (a) to delay progression of androgen-sensitive
 CC prostatic cancer cells to the androgen-independent state, in vivo or in
 CC vitro; (b) to treat prostatic cancer (after initially withdrawing
 CC androgens to induce apoptosis); and (c) to increase sensitivity of cancer
 CC cells (prostatic, renal, non-small cell lung, urothelial transitional,
 CC ovarian and some breast cancer cells) that express abnormal levels of
 CC TRPM-2 to chemotherapy or radiation. The modifications present in (I)
 CC increase stability in vivo and activity (both in vivo or in vitro) and
 CC result in a synergistic increase in effect when (I) is used with
 CC chemotherapeutic agents or other antisense oligonucleotides directed
 CC against other antiapoptotic genes. The present sequence represents a
 CC specific example of an anti-apoptotic protein TRPM-2 (testosterone-
 CC repressed prostate message-2) antisense oligonucleotide
 XX

SQ Sequence 21 BP; 6 A; 6 C; 4 G; 5 T; 0 U; 0 Other;

Query Match 90.5%; Score 19; DB 10; Length 21;
 Best Local Similarity 73.7%; Pred. No. 34;
 Matches 14; Conservative 5; Mismatches 0; Indels 0; Gaps 0;

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 Db 3 GCAGCAGAGTCTTCATCAT 21

RESULT 9

ADM83069

ID ADM83069 standard; DNA; 21 BP.

XX ADM83069;
 AC

XX 03-JUN-2004 (first entry)
 DT

XX Human TRPM-2 antisense oligonucleotide #4.
 DE

XX Testosterone-repressed prostate message-2; TRPM-2; chemo-sensitivity;
 KW radiation-sensitivity; prostate cancer; bladder cancer; ovarian cancer;
 KW lung cancer; renal cell carcinoma; RCC; antisense gene therapy; human;
 KW antisense; ss.

XX Homo sapiens.
 OS

XX Synthetic.
 OS

XX Key Location/Qualifiers
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XX 21-AUG-2003.
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XX 28-SEP-2001; 2001US-00967726.
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XX 25-FEB-2000; 2000WO-US004875.
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XX 28-SEP-2000; 2000US-0236301P.
 PR

XX 10-AUG-2001; 2001US-00913325.
 PR

XX (GLEA/) GLEAVE M.
 PA

XX (RENN/) RENNIE P S.
 PA

XX (MIYA/) MIYAKE H.
 PA

XX (NELS/) NELSON C.
 PA

XX (ZELL/) ZELLWEGER T.
 PA

XX Gleave M, Rennie PS, Miyake H, Nelson C, Zellweger T;
 PI

XX WPI; 2003-778017/73.
 DR

XX Enhancing the chemo-sensitivity or radiation-sensitivity of cancer cells
 PT that expresses testosterone-repressed prostate message-2 (TRPM-2)
 PT comprises administering a composition that inhibits expression of TRPM-2.

XX Claim 4; SEQ ID NO 4; 14pp; English.
 PS

XX The present invention provides a method for treating cancer in which
 CC cancer cells express testosterone-repressed prostate message-2 (TRPM-2).
 CC The invention is useful for enhancing the chemo-sensitivity or radiation-
 CC sensitivity of cancer cells for treating cancer such as prostate cancer,
 CC bladder cancer, ovarian cancer, lung cancer and renal cell carcinoma
 CC (RCC). The invention is also useful in antisense gene therapy. The
 CC present sequence is human testosterone-repressed prostate message-2 (TRPM-
 CC -2) antisense oligodeoxyribonucleotide (ODN).

XX Sequence 21 BP; 6 A; 6 C; 4 G; 5 T; 0 U; 0 Other;

Query Match 90.5%; Score 19; DB 11; Length 21;
 Best Local Similarity 73.7%; Pred. No. 34;
 Matches 14; Conservative 5; Mismatches 0; Indels 0; Gaps 0;

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 Db 3 GCAGCAGAGTCTTCATCAT 21

RESULT 10

ADL70464/c

ID ADL70464 standard; RNA; 21 BP.

XX

AC ADL70464;

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XX 20-MAY-2004 (first entry)
DT RNAi for human clusterin.
XX
XX RNA interference; RNAi; short interfering RNA; siRNA; human; clusterin;
DE cytosolic; neuroprotective; nootropic; gene silencing; DNA-RNA hybrid;
XX ss.
XX Homo sapiens.
OS Synthetic.
XX
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XX WO2004018676-A2.
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XX 04-MAR-2004.
XX
XX 21-AUG-2003; 2003WO-CA001277.
XX
XX 21-AUG-2002; 2002US-0405193P.
PR 03-SEP-2002; 2002US-0408152P.
PR 20-MAY-2003; 2003US-0472387P.
XX
XX (UYBR-) UNIV BRITISH COLUMBIA.
XX
XX Jansen B, Gleave ME, Signaevsky M, Beraldi E, Trougakos IP;
PI Gonos ES;
XX
XX WPI; 2004-226852/21.
XX
XX New RNA molecule less than 49 bases and having a sequence effective to
PT mediate degradation or block translation of mRNA that is the
PT transcriptional product of a target gene, useful for treating Alzheimer's
PT disease or cancer.
XX
XX Claim 4; SEQ ID NO 9; 63pp; English.
XX
XX The present sequence is the sense strand of a short interfering RNA
-XX (siRNA) targeted to human clusterin. The antisense strand is also
CC provided ADL70465. The siRNA can be used to interfere with the expression
CC of clusterin. Clusterin, also known as testosterone-repressed prostate
CC message-2 (TRPM-2) or sulfated glycoprotein-2 (SGP-2), is expressed in
CC increased amounts by prostate tumour cells following androgen withdrawal,
CC and has also been shown to be critical for neuritic toxicity in mouse
CC models of Alzheimer's disease. siRNAs of the invention can be used alone
CC or in combination with other chemotherapy or apoptosis inducing
CC treatments for the treatment of prostate cancer, sarcomas such as
CC osteosarcoma, renal cell carcinoma, breast cancer, bladder cancer, lung
CC cancer, colon cancer, ovarian cancer, anaplastic large cell lymphoma and
CC melanoma, and also for the treatment of Alzheimer's disease.
XX
XX Sequence 21 BP; 5 A; 4 C; 5 G; 2 T; 5 U; 0 Other;
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Best Local Similarity 73.7%; Pred. No. 34;
Matches 14; Conservative 5; Mismatches 0; Indels 0; Gaps 0;
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DB 19 GCAGCAGAGTCTTCATCAT 1
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ID ADL70430 standard; RNA; 21 BP.
XX
XX AC ADL70430;
XX
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DT 20-MAY-2004 (first entry)
XX RNAi for human clusterin.
XX
XX Human; clusterin; RNAi; melanoma; cytostatic; gene silencing;
DE short interfering RNA; siRNA; DNA-RNA hybrid; ss.
XX Homo sapiens.
OS Synthetic.
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XX WO2004018675-A1.
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XX 21-AUG-2003; 2003WO-CA001276.
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XX 21-AUG-2002; 2002US-0405193P.
PR 03-SEP-2002; 2002US-0408152P.
PR 02-DEC-2002; 2002US-0319748P.
PR 20-MAY-2003; 2003US-0472387P.
XX
XX (UYBR-) UNIV BRITISH COLUMBIA.
XX
XX (GLEA/) GLEAVE M E.
XX
XX Jansen B;
XX
XX WPI; 2004-226851/21.
XX
XX Treating melanoma in a mammalian subject comprises administering to the
PT subject a therapeutic agent effective to reduce the effective amount of
PT clusterin in the melanoma cells.
XX
XX Claim 20; SEQ ID NO 28; 32pp; English.
XX
XX The present sequence is that of a short interfering RNA (siRNA) molecule
CC targeted to human clusterin ADL70403. The invention relates to the
CC treatment of melanoma through reduction in the effective amount of
CC clusterin. The therapeutic agent may be an antisense oligonucleotide
CC ADL70404-ADL70421 or short interfering RNA (siRNA) ADL70422-ADL70445
CC targeted to clusterin. The siRNAs molecules direct cleavage of clusterin
CC mRNA. A method for regulating expression of bcl-xL in a subject or cell
CC line comprises administering an agent effective to modulate the amount of
CC clusterin expression. In clusterin-expressing cells, expression of bcl-xL
CC is down-regulated when the effective amount of clusterin is reduced. Such
CC inhibition is significant because bcl-xL is known to act as an inhibitor
CC of apoptosis.
XX
XX Sequence 21 BP; 5 A; 4 C; 5 G; 2 T; 5 U; 0 Other;
SQ
Query Match 90.5%; Score 19; DB 12; Length 21;
Best Local Similarity 73.7%; Pred. No. 34;
Matches 14; Conservative 5; Mismatches 0; Indels 0; Gaps 0;
QY 1 GCAGCAGAGUCUUCAUCAU 19
DB 19 GCAGCAGAGTCTTCATCAT 1
RESULT 12
ADL70406
ID ADL70406 standard; DNA; 21 BP.
XX
XX AC ADL70406;
XX
XX 20-MAY-2004 (first entry)
XX
XX Antisense oligonucleotide to human clusterin.
DE
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XX Human; clusterin; antisense; melanoma; cytostatic; gene silencing; ss.
XX Homo sapiens.
OS Synthetic.
XX Key Location/Qualifiers
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XX WO2004018675-A1.
XX 04-MAR-2004.
XX 21-AUG-2003; 2003WO-CA001276.
XX 21-AUG-2002; 2002US-0405193P.
XX 03-SEP-2002; 2002US-0408152P.
XX 02-DEC-2002; 2002US-0319748P.
XX 20-MAY-2003; 2003US-0472387P.
XX (UYBR-) UNIV BRITISH COLUMBIA.
XX (GLEA/) GLEAVE M E.
XX Jansen B;
XX WPI; 2004-226851/21.
XX Treating melanoma in a mammalian subject comprises administering to the
XX subject a therapeutic agent effective to reduce the effective amount of
XX clusterin in the melanoma cells.
XX Claim 7; SEQ ID NO 4; 32pp; English.
XX The present sequence is that of an antisense oligonucleotide targeted to
XX human clusterin ADL70403. The invention relates to the treatment of
XX melanoma through reduction in the effective amount of clusterin. The
XX therapeutic agent may be an antisense oligonucleotide ADL70404-ADL70421
XX or short interfering RNA (siRNA) ADL70422-ADL70445 targeted to clusterin.
XX The antisense oligonucleotides are complementary to a region of the
XX clusterin mRNA spanning either the translation initiation site or the
XX termination site. They may be modified to increase stability in vivo,
XX e.g. they may be employed as phosphorothioate derivatives and may have 2'
XX -O-(2-methoxyethyl) (MOE) modifications in the 5' and 3' 'wings'. The
XX present antisense oligonucleotide is particularly preferred. It is
XX targeted to the translation initiation codon and next 6 codons of the
XX human clusterin sequence. It has a phosphorothioate backbone throughout
XX and MOE wings, the remaining nucleotides being 2'-deoxynucleotides. In an
XX example from the invention, this antisense oligonucleotide provided a
XX dose-dependent down-regulation of clusterin in human melanoma cells,
XX leading to an increase in apoptotic cell death. In one melanoma cell line
XX (607B) this alone was sufficient to lead to complete cell death. In
XX another melanoma cell line, the surviving cells showed increased
XX sensitivity to subsequent treatment with cisplatin. A claimed method for
XX regulating expression of bcl-xL in a subject or cell line comprises
XX administering an agent effective to modulate the amount of clusterin
XX expression. In clusterin-expressing cells, expression of bcl-xL is down-
XX regulated when the effective amount of clusterin is reduced. Such
XX inhibition is significant because bcl-xL is known to act as an inhibitor
XX of apoptosis.
XX Sequence 21 BP; 6 A; 6 C; 4 G; 5 T; 0 U; 0 Other;

Query Match 90.5%; Score 19; DB 12; Length 21;
Best Local Similarity 73.7%; Pred. No. 34;
Matches 14; Conservative 5; Mismatches 0; Indels 0; Gaps 0;
QY 1 GCAGCAGAGUCUUCAU 19
DB 3 GCAGCAGAGTCTTCATCAT 21
RESULT 13
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ID ADL70521 standard; cDNA; 23 BP.
XX AC ADL70521;
XX 20-MAY-2004 (first entry)
XX Human clusterin target for RNAi.
XX RNA interference; RNAi; short interfering RNA; siRNA; human; clusterin;
XX cytostatic; neuroprotective; nootropic; gene silencing; DNA-RNA hybrid;
XX ss.
XX Homo sapiens.
OS Synthetic.
XX WO2004018676-A2.
XX 04-MAR-2004.
XX 21-AUG-2003; 2003WO-CA001277.
XX 21-AUG-2002; 2002US-0405193P.
XX 03-SEP-2002; 2002US-0408152P.
XX 20-MAY-2003; 2003US-0472387P.
XX (UYBR-) UNIV BRITISH COLUMBIA.
XX Jansen B, Gleave ME, Signaevsky M, Beraldi E, Trougakos IP;
XX Gonos ES;
XX WPI; 2004-226852/21.
XX New RNA molecule less than 49 bases and having a sequence effective to
XX mediate degradation or block translation of mRNA that is the
XX transcriptional product of a target gene, useful for treating Alzheimer's
XX disease or cancer.
XX Example 6; SEQ ID NO 66; 63pp; English.
XX The present sequence is a human clusterin cDNA target for a double-
XX stranded short interfering RNA (siRNA) of the invention to demonstrate
XX ADL70523. It was used in an example from the invention to demonstrate
XX clusterin gene silencing in PC-3 prostate cancer cells. Clusterin, also
XX known as testosterone-repressed prostate message-2 (TRPM-2) or sulfated
XX glycoprotein-2 (SGP-2), is expressed in increased amounts by prostate
XX tumour cells following androgen withdrawal, and has also been shown to be
XX critical for neuritic toxicity in mouse models of Alzheimer's disease.
XX siRNAs of the invention can be used alone or in combination with other
XX chemotherapy or apoptosis inducing treatments for the treatment of
XX prostate cancer, sarcomas such as osteosarcoma, renal cell carcinoma,
XX breast cancer, bladder cancer, lung cancer, colon cancer, ovarian cancer,
XX anaplastic large cell lymphoma and melanoma, and also for the treatment
XX of Alzheimer's disease.
XX Sequence 23 BP; 5 A; 5 C; 7 G; 6 T; 0 U; 0 Other;
QY 1 GCAGCAGAGUCUUCAU 19
DB 3 GCAGCAGAGTCTTCATCAT 21
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Best Local Similarity 73.7%; Pred. No. 34;
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C 5	19	90.5	19	6	CO786653	Sequence	
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C 7	19	90.5	21	6	CO786121	Sequence	
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C 9	19	90.5	21	6	CO786639	Sequence	
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C 12	19	90.5	195	6	AR302784	Sequence	
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C 14	19	90.5	270	6	CO688092	Sequence	
C 15	19	90.5	275	6	AX895152	Sequence	
C 16	19	90.5	275	6	ED030685	Sequence	
C 17	19	90.5	306	6	CO684015	Sequence	
C 18	19	90.5	396	6	AR391192	Sequence	
C 19	19	90.5	396	6	AR392897	Sequence	

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other sequences; artificial sequences.
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REFERENCE
AUTHORS Jansen,B.
TITLE Treatment of melanoma by reduction in clusterin levels
JOURNAL Patent: WO 2004018675-A 29 04-MAR-2004;
The University of British Columbia (CA); Gleave, Martin E. (CA)
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LOCUS
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ACCESSION CQ786179
VERSION CQ786179.1 GI:45721282
KEYWORDS
SOURCE synthetic construct
ORGANISM other sequences; artificial sequences.
REFERENCE
AUTHORS Jansen,B., Gleave,M.E., Signaevsky,M., Beraldi,E., Trougakos,I. and
Gonos,E.
TITLE Rnai probes targeting cancer-related proteins
JOURNAL Patent: WO 2004018676-A 67 04-MAR-2004;
The University of British Columbia (CA)
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LOCUS
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ACCESSION CQ786180
VERSION CQ786180.1 GI:45721283
KEYWORDS
SOURCE synthetic construct
ORGANISM other sequences; artificial sequences.
REFERENCE
AUTHORS Jansen,B., Gleave,M.E., Signaevsky,M., Beraldi,E., Trougakos,I. and
Gonos,E.
TITLE Rnai probes targeting cancer-related proteins
JOURNAL Patent: WO 2004018676-A 68 04-MAR-2004;
The University of British Columbia (CA)
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CQ786653/c CQ786653 19 bp RNA linear PAT 24-MAR-2004
LOCUS
DEFINITION Sequence 42 from Patent WO2004018675.
ACCESSION CQ786653
VERSION CQ786653.1 GI:45721673
KEYWORDS
SOURCE synthetic construct
ORGANISM other sequences; artificial sequences.
REFERENCE
AUTHORS Jansen,B.
TITLE Treatment of melanoma by reduction in clusterin levels
JOURNAL Patent: WO 2004018675-A 42 04-MAR-2004;
The University of British Columbia (CA); Gleave, Martin E. (CA)
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Matches 14; Conservative 5; Mismatches 0; Indels 0; Gaps 0;
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Db 1 GCAGCAGAGTCTTCATCAT 19
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CQ786654 CQ786654 19 bp RNA linear PAT 24-MAR-2004
LOCUS
DEFINITION Sequence 43 from Patent WO2004018675.
ACCESSION CQ786654
VERSION CQ786654.1 GI:45721674
KEYWORDS
SOURCE synthetic construct
ORGANISM other sequences; artificial sequences.
REFERENCE
AUTHORS Jansen,B.
TITLE Treatment of melanoma by reduction in clusterin levels
JOURNAL Patent: WO 2004018675-A 43 04-MAR-2004;
The University of British Columbia (CA); Gleave, Martin E. (CA)
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/note="RNAi for human clusterin"
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Best Local Similarity 73.7%; Pred. No. 3.1e+02;
Matches 14; Conservative 5; Mismatches 0; Indels 0; Gaps 0;
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Db 19 GCAGCAGAGTCTTCATCAT 1

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RESULT 9

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Job time : 1718 secs



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GenCore version 5.1.6
Copyright (c) 1993 - 2005 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: May 23, 2005, 23:16:37 ; Search time 341 Seconds
(without alignments)
377.650 Million cell updates/sec

Title: US-10-646-436-10

Perfect score: 21

Sequence: 1 gcagcagagucuccauctt 21

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Total number of hits satisfying chosen parameters: 11390874

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SUMMARIES

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3	19	90.5	19	17	US-10-646-391A-42
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6	19	90.5	19	17	US-10-646-436-68
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8	19	90.5	21	10	US-09-967-736A-4
9	19	90.5	21	16	US-10-080-794-4
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11	19	90.5	21	17	US-10-646-391A-28

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Sequence 37963, A
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Sequence 33018, A
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Sequence 28941, A
Sequence 28941, A
Sequence 2963, Ap
Sequence 15, Appl
Sequence 15, Appl
Sequence 15, Appl
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Sequence 32172, A
Sequence 2524, Ap
Sequence 32177, A
Sequence 12511, A
Sequence 17455, A
Sequence 31156, A
Sequence 1985, Ap
Sequence 91, Appl
Sequence 93, Appl
Sequence 101, App
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Sequence 29, Appl
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ALIGNMENTS

RESULT 1
US-10-646-391A-29
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; Publication NO. US20040082534A1
; GENERAL INFORMATION:
; APPLICANT: Jansen, Martin
; TITLE OF INVENTION: Treatment of Melanoma by Reduction in Clusterin Levels
; FILE REFERENCE: UBC.P-035
; CURRENT APPLICATION NUMBER: US/10/646,391A
; PRIOR FILING DATE: 2003-08-21
; PRIOR APPLICATION NUMBER: US 60/405,193
; PRIOR FILING DATE: 2002-08-21
; PRIOR APPLICATION NUMBER: US 60/319,748
; PRIOR FILING DATE: 2002-12-02
; PRIOR APPLICATION NUMBER: US 60/408,152
; PRIOR FILING DATE: 2002-09-03
; PRIOR APPLICATION NUMBER: US 60/473,387
; PRIOR FILING DATE: 2003-05-20
; NUMBER OF SEQ ID NOS: 43
; SOFTWARE: Patentin version 3.2
; SEQ ID NO 29
; LENGTH: 21
; TYPE: DNA
; ORGANISM: artificial
; FEATURE:
; OTHER INFORMATION: RNAi for human clusterin
US-10-646-391A-29

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; Publication No. US20040096882A1
; GENERAL INFORMATION:
; APPLICANT: Jansen, Burkhard
; APPLICANT: Gleave, Martin
; APPLICANT: Signaevsky, Maxim
; APPLICANT: Beraldi, Eliana
; APPLICANT: Trougakos, Ioannis
; APPLICANT: Gonos, Efsthathios
; TITLE OF INVENTION: RNAi Probes Targeting Cancer-Related Proteins
; FILE REFERENCE: UBC.P-030
; CURRENT APPLICATION NUMBER: US/10/646,436
; CURRENT FILING DATE: 2003-08-21
; PRIOR APPLICATION NUMBER: US 60/405,193
; PRIOR FILING DATE: 2002-08-21
; PRIOR APPLICATION NUMBER: US 60/408,152
; PRIOR FILING DATE: 2002-09-03
; PRIOR APPLICATION NUMBER: US 60/473,387
; PRIOR FILING DATE: 2003-05-20
; NUMBER OF SEQ ID NOS: 68
; SOFTWARE: PatentIn version 3.2
; SEQ ID NO 10
; LENGTH: 21
; TYPE: DNA
; ORGANISM: artificial
; FEATURE:
; OTHER INFORMATION: RNAi for human clusterin
US-10-646-436-10

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Qy      1 GCAGCAGAGUCUUCAUCAUTT 21
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; Publication No. US20040082534A1
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; APPLICANT: Jansen, Burkhard
; APPLICANT: Gleave, Martin
; TITLE OF INVENTION: Treatment of Melanoma by Reduction in Clusterin Levels
; FILE REFERENCE: UBC.P-035
; CURRENT APPLICATION NUMBER: US/10/646,391A
; CURRENT FILING DATE: 2003-08-21
; PRIOR APPLICATION NUMBER: US 60/405,193
; PRIOR FILING DATE: 2002-08-21
; PRIOR APPLICATION NUMBER: US 60/319,748
; PRIOR FILING DATE: 2002-12-02
; PRIOR APPLICATION NUMBER: US 60/408,152
; PRIOR FILING DATE: 2002-09-03
; PRIOR APPLICATION NUMBER: US 60/473,387
; PRIOR FILING DATE: 2003-05-20
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; TYPE: RNA
; ORGANISM: artificial
; FEATURE:
; OTHER INFORMATION: RNAi for human clusterin
US-10-646-391A-42

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; Sequence 67, Application US/10646436
; Publication No. US20040096882A1
; GENERAL INFORMATION:
; APPLICANT: Jansen, Burkhard
; APPLICANT: Gleave, Martin
; APPLICANT: Signaevsky, Maxim
; APPLICANT: Beraldi, Eliana
; APPLICANT: Trougakos, Ioannis
; APPLICANT: Gonos, Efsthathios
; TITLE OF INVENTION: RNAi Probes Targeting Cancer-Related Proteins
; FILE REFERENCE: UBC.P-030
; CURRENT APPLICATION NUMBER: US/10/646,436
; CURRENT FILING DATE: 2003-08-21
; PRIOR APPLICATION NUMBER: US 60/405,193
; PRIOR FILING DATE: 2002-08-21
; PRIOR APPLICATION NUMBER: US 60/408,152
; PRIOR FILING DATE: 2002-09-03
; PRIOR APPLICATION NUMBER: US 60/473,387
; PRIOR FILING DATE: 2003-05-20
; NUMBER OF SEQ ID NOS: 68
; SOFTWARE: PatentIn version 3.2
; SEQ ID NO 67
; LENGTH: 19
; TYPE: RNA
; OTHER INFORMATION: RNAi for human clusterin
US-10-646-436-67/c
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; ORGANISM: artificial
; FEATURE:
; OTHER INFORMATION: RNAi for human clusterin
US-10-646-436-67

Query Match 90.5%; Score 19; DB 17; Length 19;
Best Local Similarity 73.7%; Pred. No. 26;
Matches 14; Conservative 5; Mismatches 0; Indels 0; Gaps 0;

QY 1 GCAGCAGAGUCUUCAU 19
Db 19 GCAGCAGAGTCCTTCATCAT 1

RESULT 6
US-10-646-436-68
; Sequence 68, Application US/10646436
; Publication No. US20040096882A1
; GENERAL INFORMATION:
; APPLICANT: Jansen, Burkhard
; APPLICANT: Gleave, Martin
; APPLICANT: Sigmaevsky, Maxim
; APPLICANT: Beraldi, Eliana
; APPLICANT: Trougakos, Ioannis
; APPLICANT: Gonos, Efstathios
; TITLE OF INVENTION: RNAi Probes Targeting Cancer-Related Proteins
; FILE REFERENCE: UBC.P-030
; CURRENT APPLICATION NUMBER: US/10/646,436
; CURRENT FILING DATE: 2003-08-21
; PRIOR APPLICATION NUMBER: US 60/405,193
; PRIOR FILING DATE: 2002-08-21
; PRIOR APPLICATION NUMBER: US 60/408,152
; PRIOR FILING DATE: 2002-09-03
; PRIOR APPLICATION NUMBER: US 60/473,387
; PRIOR FILING DATE: 2003-05-20
; NUMBER OF SEQ ID NOS: 68
; SOFTWARE: PatentIn version 3.2
; SEQ ID NO 68
; LENGTH: 19
; TYPE: RNA
; ORGANISM: artificial
; FEATURE:
; OTHER INFORMATION: RNAi fo rhuman clusterin
US-10-646-436-68

Query Match 90.5%; Score 19; DB 17; Length 19;
Best Local Similarity 100.0%; Pred. No. 26;
Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GCAGCAGAGUCUUCAU 19
Db 1 GCAGCAGAGUCUUCAU 19

RESULT 7
US-09-944-326-4
; Sequence 4, Application US/09944326
; Patent No. US20020128220A1
; GENERAL INFORMATION:
; APPLICANT: Gleave, Martin
; APPLICANT: Rennie, Paul S.
; APPLICANT: Miyake, Hideaki
; APPLICANT: Nelson, Colleen
; TITLE OF INVENTION: TRPM-2 ANTISENSE THERAPY
; FILE REFERENCE: UBC.P-020-2
; CURRENT APPLICATION NUMBER: US/09/944,326
; CURRENT FILING DATE: 2001-08-30
; PRIOR APPLICATION NUMBER: 60/121,726
; PRIOR FILING DATE: 1999-02-26
; PRIOR APPLICATION NUMBER: 09/913,325
; PRIOR FILING DATE: 2001-08-10
; NUMBER OF SEQ ID NOS: 14
; SOFTWARE: PatentIn Ver. 2.1

; SEQ ID NO 4
; LENGTH: 21
; TYPE: DNA
; ORGANISM: HUMAN
; FEATURE:
; OTHER INFORMATION: antisense TRPM-2 ODN
US-09-944-326-4

Query Match 90.5%; Score 19; DB 9; Length 21;
Best Local Similarity 73.7%; Pred. No. 27;
Matches 14; Conservative 5; Mismatches 0; Indels 0; Gaps 0;

QY 1 GCAGCAGAGUCUUCAU 19
Db 3 GCAGCAGAGTCCTTCATCAT 21

RESULT 8
US-09-967-726A-4
; Sequence 4, Application US/09967726A
; Publication No. US20030158130A1
; GENERAL INFORMATION:
; APPLICANT: Gleave, Martin
; APPLICANT: Rennie, Paul S.
; APPLICANT: Miyake, Hideaki
; APPLICANT: Nelson, Colleen
; APPLICANT: Zellweger, Tobias
; TITLE OF INVENTION: Chemo- and Radiation-Sensitization of Cancer by Antisense TRPM-2
; TITLE OF INVENTION: Oligonucleotides
; FILE REFERENCE: UBC.P-022
; CURRENT APPLICATION NUMBER: US/09/967,726A
; CURRENT FILING DATE: 2001-09-28
; NUMBER OF SEQ ID NOS: 15
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 4
; LENGTH: 21
; TYPE: DNA
; ORGANISM: human
; ORGANISM: human
US-09-967-726A-4

Query Match 90.5%; Score 19; DB 10; Length 21;
Best Local Similarity 73.7%; Pred. No. 27;
Matches 14; Conservative 5; Mismatches 0; Indels 0; Gaps 0;

QY 1 GCAGCAGAGUCUUCAU 19
Db 3 GCAGCAGAGTCCTTCATCAT 21

RESULT 9
US-10-080-794-4
; Sequence 4, Application US/10080794
; Publication No. US20030166591A1
; GENERAL INFORMATION:
; APPLICANT: Gleave, Martin
; APPLICANT: Rennie, Paul S.
; APPLICANT: Miyake, Hideaki
; APPLICANT: Nelson, Colleen
; APPLICANT: Monia, Brett P.
; TITLE OF INVENTION: TRPM-2 ANTISENSE THERAPY USING AN OLIGONUCLEOTIDE
; FILE REFERENCE: UBC.P-020-3
; CURRENT APPLICATION NUMBER: US/10/080,794
; CURRENT FILING DATE: 2002-02-22
; PRIOR APPLICATION NUMBER: 60/121,726
; PRIOR FILING DATE: 1999-02-26
; PRIOR APPLICATION NUMBER: 09/913,325
; PRIOR FILING DATE: 2001-08-10
; PRIOR APPLICATION NUMBER: 09/944,326
; PRIOR FILING DATE: 2001-08-30
; NUMBER OF SEQ ID NOS: 19
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 4

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; LENGTH: 21
; TYPE: DNA
; ORGANISM: HUMAN
; FEATURE:
; OTHER INFORMATION: antisense TRPM-2 ODN
US-10-080-794-4

Query Match          90.5%; Score 19; DB 16; Length 21;
Best Local Similarity 73.7%; Pred. No. 27;
Matches 14; Conservative 5; Mismatches 0; Indels 0; Gaps 0;

Qy 1 GCAGCAGAGUCUUCAUCAU 19
Db 3 GCAGCAGAGTCTTCATCAT 21

RESULT 10
US-10-646-391A-4
; Sequence 4, Application US/10646391A
; Publication No. US20040082534A1
; GENERAL INFORMATION:
; APPLICANT: Jansen, Burkhard
; TITLE OF INVENTION: Treatment of Melanoma by Reduction in Clusterin Levels
; FILE REFERENCE: UBC P-035
; CURRENT APPLICATION NUMBER: US/10/646,391A
; CURRENT FILING DATE: 2003-08-21
; PRIOR APPLICATION NUMBER: US 60/405,193
; PRIOR FILING DATE: 2002-08-21
; PRIOR APPLICATION NUMBER: US 60/319,748
; PRIOR FILING DATE: 2002-12-02
; PRIOR APPLICATION NUMBER: US 60/408,152
; PRIOR FILING DATE: 2002-09-03
; PRIOR APPLICATION NUMBER: US 60/473,387
; PRIOR FILING DATE: 2003-05-20
; NUMBER OF SEQ ID NOS: 43
; SOFTWARE: PatentIn version 3.2
; SEQ ID NO 4
; LENGTH: 21
; TYPE: DNA
; ORGANISM: human
US-10-646-391A-4

Query Match          90.5%; Score 19; DB 17; Length 21;
Best Local Similarity 73.7%; Pred. No. 27;
Matches 14; Conservative 5; Mismatches 0; Indels 0; Gaps 0;

Qy 1 GCAGCAGAGUCUUCAUCAU 19
Db 3 GCAGCAGAGTCTTCATCAT 21

RESULT 11
US-10-646-391A-28/c
; Sequence 28, Application US/10646391A
; Publication No. US20040082534A1
; GENERAL INFORMATION:
; APPLICANT: Jansen, Burkhard
; TITLE OF INVENTION: Treatment of Melanoma by Reduction in Clusterin Levels
; FILE REFERENCE: UBC P-035
; CURRENT APPLICATION NUMBER: US/10/646,391A
; CURRENT FILING DATE: 2003-08-21
; PRIOR APPLICATION NUMBER: US 60/405,193
; PRIOR FILING DATE: 2002-08-21
; PRIOR APPLICATION NUMBER: US 60/319,748
; PRIOR FILING DATE: 2002-12-02
; PRIOR APPLICATION NUMBER: US 60/408,152
; PRIOR FILING DATE: 2002-09-03
; PRIOR APPLICATION NUMBER: US 60/473,387
; PRIOR FILING DATE: 2003-05-20
; NUMBER OF SEQ ID NOS: 43
; SOFTWARE: PatentIn version 3.2

; SEQ ID NO 28
; LENGTH: 21
; TYPE: DNA
; ORGANISM: artificial
; FEATURE:
; OTHER INFORMATION: RNAi for human clusterin
US-10-646-391A-28

Query Match          90.5%; Score 19; DB 17; Length 21;
Best Local Similarity 73.7%; Pred. No. 27;
Matches 14; Conservative 5; Mismatches 0; Indels 0; Gaps 0;

Qy 1 GCAGCAGAGUCUUCAUCAU 19
Db 19 GCAGCAGAGTCTTCATCAT 1

RESULT 12
US-10-646-436-9/c
; Sequence 9, Application US/10646436
; Publication No. US2004009882A1
; GENERAL INFORMATION:
; APPLICANT: Jansen, Burkhard
; APPLICANT: Gleave, Martin
; APPLICANT: Signaevsky, Maxim
; APPLICANT: Beraldi, Eliana
; APPLICANT: Trougakos, Ioannis
; APPLICANT: Gonos, Efstathios
; TITLE OF INVENTION: RNAi Probes Targeting Cancer-Related Proteins
; FILE REFERENCE: UBC P-030
; CURRENT APPLICATION NUMBER: US/10/646,436
; CURRENT FILING DATE: 2003-08-21
; PRIOR APPLICATION NUMBER: US 60/405,193
; PRIOR FILING DATE: 2002-08-21
; PRIOR APPLICATION NUMBER: US 60/408,152
; PRIOR FILING DATE: 2002-09-03
; PRIOR APPLICATION NUMBER: US 60/473,387
; PRIOR FILING DATE: 2003-05-20
; NUMBER OF SEQ ID NOS: 68
; SOFTWARE: PatentIn version 3.2
; SEQ ID NO 9
; LENGTH: 21
; TYPE: DNA
; ORGANISM: artificial
; FEATURE:
; OTHER INFORMATION: RNAi for human clusterin
US-10-646-436-9

Query Match          90.5%; Score 19; DB 17; Length 21;
Best Local Similarity 73.7%; Pred. No. 27;
Matches 14; Conservative 5; Mismatches 0; Indels 0; Gaps 0;

Qy 1 GCAGCAGAGUCUUCAUCAU 19
Db 19 GCAGCAGAGTCTTCATCAT 1

RESULT 13
US-10-828-394-5
; Sequence 5, Application US/10828394
; Publication No. US20040220131A1
; GENERAL INFORMATION:
; APPLICANT: Jackson, John
; APPLICANT: Burt, Helen
; APPLICANT: Springate, Christopher
; APPLICANT: Gleave, Martin
; TITLE OF INVENTION: Method for Treatment of Cancerous Angiogenic Disorders
; FILE REFERENCE: UBC P-033
; CURRENT APPLICATION NUMBER: US/10/828,394
; CURRENT FILING DATE: 2004-04-19
; PRIOR APPLICATION NUMBER: US 60/464,159
; PRIOR FILING DATE: 2003-04-18
; NUMBER OF SEQ ID NOS: 23
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RESULT 15
US-10-646-436-66/c
; Sequence 66, Application US/10646436
: Publication No. US20040096882A1

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